ALKA-SELTZER GOLD- alka-seltzer gold tablet, effervescent Bayer HealthCare LLC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Alka-Seltzer Gold

Drug Facts

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Active Ingredients

Anhydrous citric acid 1000 mg	Antacid
Potassium bicarbonate 344 mg	Antacid
Sodium bicarbonate (heat-treated) 1050 mg	Antacid

Purpose

Antacid

Uses for the relief of

• heartburn • acid indigestion • sour stomach

Do Not Use

Do not use if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

- kidney disease
- a potassium or sodium-restricted diet

Ask a doctor or pharmacist before use if you are

Ask a doctor or pharmacist before use if you are presently taking a prescription drug. Antacids may interact with certain prescription drugs.

When using this product

When using this product do not exceed recommended dosage

Stop use and ask a doctor if

Stop use and ask a doctor if you have taken the maximum dose for 2 weeks

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children

Keep out of reach of children.

Directions

• fully dissolve tablets in 4 ounces of water before taking

adults and children 12 years	2 tablets every 4 hours as needed, or	do not exceed8 tablets in
and over	asdirected by a doctor	24 hours
adults 60 years and over	2 tablets every 4 hours as needed, or as directed by a doctor	do not exceed 6 tablets in 24 hours
children under 12 years	1 tablet every 4 hours as needed, or as directed by a doctor	do not exceed 4 tablets in 24 hours

Other information

• each tablet contains: potassium 135 mg

• each tablet contains: sodium 309 mg

- store at room temperature. Avoid excessive heat.
- this product does not contain aspirin
- Alka-Seltzer Gold in water contains principally the

antacids sodium citrate and potassium citrate

Inactive ingredients

Inactive ingredients magnesium stearate, mannitol

Questions or comments

Questions or comments?1-800-986-0369 (Mon – Fri 9AM – 5PM EST)

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Allergy alert: Aspirin may cause a severe allergic reaction which may include:

· hives · facial swelling · asthma (wheezing) · shock

Stomach bleeding warning: This product contains an NSAID, which may cause severe stomach bleeding. The chance is

higher if you

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription

NSAIDs (aspirin, ibuprofen, naproxen, or others)

• have 3 or more alcoholic drinks every day while using this

product

• take more or for a longer time than directed



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LOT & EXP



ALKA-SELTZER GOLD

alka-seltzer gold tablet, effervescent

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0280-4100
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968 PHW8 QP) (ANHYDROUS CITRIC ACID - UNII:XF417D3PSL)	ANHYDROUS CITRIC ACID	1000 mg	
POTASSIUM BICARBONATE (UNII: HM5Z15LEBN) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CATION	344 mg	
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM BICARBONATE	1050 mg	

Inactive Ingredients		
Ingredient Name	Strength	
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MANNITOL (UNII: 3OWL53L36A)		

Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	25mm
Flavor		Imprint Code	
Contains			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:0280-4100-63	18 in 1 CARTON	11/19/2014		

1	2 in 1 POUCH; Type 0: Not a Combination Product		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	11/19/2014	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 5/2017 Bayer HealthCare LLC.